1. HOME DISTRICT BOS	2. REPORTING UNIT SYMBOL NOL				3. CENTRAL FILE NO. 1234567				4. J.D./T.A.	5. COU		6. DATE 8-2-99
7. PRODUCT CODE 45AF-19	8. OPERATION 13				9. PROGRAM ASSIGNMENT CODE 09001							10. HOURS 1/2
11. IDENTIFICATION (Quote pertinent labeling including Establishment name and address)  "NO CLUMP" BRAND ANTI-CAKING AGENT CLUMPLESS CORP. 3214 WHARF AVE.												
WALTHAM, MA 02154												
12. MANUFACTURER CONTROL CODES (Labels, packaging and shipping containers) BAGS CODED:					13. AMOUNT ON HAI 1200/100# BAG				SS	14. DATE LOT RECEIVED 7-15-99		
"AC 123171"	'AC 123171"				15. ESTIMATED VAL \$ 24,000.00				UE	E 16. SAMPLE NO(s). NONE		
17. DEALER (Name, street address, city, state, and ZIP code CREOLE INDUSTRIES 239 CANAL ST. NEW ORLEANS, LA 70130				ŕ		18. DISTRIBUTOR SHIPPER  XX MANUFACTURER OTHER  (Name, street address, city, state, ZIP code, and telephone)  CLUMPLESS CORP.  3214 WHARF AVE.  WALTHAM, MA 02154  (617) 765-4321						
19. ESTABLISHMENT TYPE(s) INDU			JSTF 3	RY C	CODE 5	6		TABLISHMENT <i>VOLUME)</i>	SIZE	OI	FORMATION BTAINED BY	
a. Manufacutur	·or	45		$\rightarrow$							(C	Check one) MAIL
b.	GI					+ +						TELEPHONE
C.											XXX	VISIT
22. REMARKS												
23. REPORT PREPARED BY (Type or print name and title)						4. EMF	PLOYE	EE NO.	25. PC 2	26. SIGNATURWE Sidney H. Rogers		
Sidney H. Rogers, Investigator					$\perp$	073				· · ·		
27. REPORTING UNIT ACTION  XX REFERRED TO HOME DISTRICT  COLLECT OFFICIAL SAMPLE  ROUTINE FOLLOW-UP									28. NAME OF REVIEWING OFFICIAL (Type or print) Harry Abelman			
REFERRED TO STATE OR INSPECT OTHER FEDERAL AUTHORITIES						T NO A	) ACTION		Supervisory			D. DATE REVIEWED

FORM FDA 457 (5/90)

PREVIOUS EDITION MAY BE USED

PRODUCT/ESTABLISHMENT

**SURVEILLANCE REPORT** 

	SUSPECTED VIOLATION	IS (Ch	eck appropriate box)							
		HEALTH	,							
	Dangerous under any condition of use: 502(j).		Inadequate directions for use: 502(f)(1).							
	Dangerous when sold indiscriminately: 502(f).		Failure to bear list of active ingredients: 502(e).							
DRUGS – DEVICES	Dangerous on account of excessive dosage: 502(j).		Possible variation from professed standard: 501(b), (c),							
	Dangerous because of inadequate warnings: 502(f)(2).		Vitamin preparations – possible variation from professed standard: 501(b), (c), (d).							
	Drugs dangerous on account of impurities: 501(a)(2), (3), 502(j).		Extravagant therapeutic claims: 502(a) <sup>1</sup>							
	If descriptive or promotional material employed in sale of product bears or contains extravagant therapeutic claims, indicate (in RE on front or in separate memo) source, how received, and how employed in sale of product. See Section 201(m), Labeling: 301(b), 3 Prohibited Acts.									
	☐ HYGIENIC									
	EC	CONOMIC								
	Deceptive packaged: 502(I).		Suspect short weight or volume: 502(b)							
	□ NEW DRUG									
		HEALTH								
FOODS	Presence of poisons: 402(a)(1), (2).		Therapeutic claims for food: Subject to 502.							
	Dangerous and non-nutritive substances (confectionery): 402(d)		Vitamin claims: 403(a), (j). May also be subject to 502.							
	Poisonous containers: 402(a)(6).		Special dietary foods: 403(j)							
		YGIENIC	T =							
	Stored under insanitary conditions: 402(a)(4).		Suspected filth or decomposition: 402(a)(3).							
		CONOMIC								
	Deceptive packaging: 403(d).		Failure to declare mandatory statements: nonstandardi foods: 403(e), (f), (l), (k).							
	Short weight or volume: 403(e)(2).		Standardized foods, misbranding or nonconformity: 403 (h).							
	Misrepresentation in labeling: 403(a). See 201(m).	X	New Product, New Manufacturer							
METICS	Dangerous cosmetics: 601(a).		Adulteration: 601.							
COSME	Misbranding: 602									
	EXPLAIN									
~										
отнек										
ОТ										

FORM FDA 457 (5/90) (BACK)